Mother's or Infant's Nan	Mother's or Infant's Name (in case of page separation):																													
HOSPITAL INFORMATION													ON														I			
														R'S MEDICAL RECORD H HOSPITAL:																
																RD														
												INFANT'S MEDICAL RECORD AT TRANSFER HOSPITAL:																		
42. DATE OF TR. TRANSFER H																				/										
									Р	HYS	ICIA	N INF	OR	MAT	ION															
N. Carterina de la carterina de													MC	TELEPHONE NUMBER OF MOTHER'S PHYSICIAN: (with area code)																
45. NAME AND ADDRESS OF INFANT'S PHYSICIAN:												46. TELEPHONE NUMBER OF INFANT'S PHYSICIAN:												7-						
(with area code) BIRTH DEFECT DIAGNOSIS																														
A. CHROMOSOMAL ABNORMALITIES Do not report: a) Heterochromatin Variants; b) Satellite/Stalk Variants of Chromosomes 13,14,15,21,22; c) Inv(2) (p11;q13); d) Inv (9) (p11;q12 or q13); e) Familial Y Variants; or d) Pse																														
Do not report: a) Heterochron 47. CYTOGENETIC	matin Va	ariants; t	o) Sate	ellite/St	alk Va	riants	of Chr	omoso	omes 1	3,14,1	15,21,2	22; c) I	nv(2)	(p11;c	13); d) Inv (9) (p11	;q12 o	r q13);	e) Fam	ilial Y	' Varia	ints; c	or d) Ps T	se	1	1		_	
DIASNOSIS (ISCN																														
Short Form)																														
48. INHERITANCE OF	49. CYTOGENETIC LAB SPECIMEN NUMBER:															51. S	PECIN	MEN TY	PE:		LIVEBORN BLOOD									
STRUCTURAL REARRANGEMENTS:																	C FLUI		LIVEBORN TISSI								OD.			
DE NOVO													CHORIONIC VILLU PERCUTANEOUS I									· · ·								
PATERNAL 50. LAB NAME: MATERNAL														ABORTUS SPECIMEN OTHER Spec																
UNKNOWN															STILLBORN TISSUE/BLOOD															
52. SAMPLING DATECheck if final result date used 53. GESTATIONAL AGE AT TIME OF SAMPLING: (Prenatal: Abortus, Fetal												We		54. METHOD USED TO DETERMINE GESTATIONAL AGE AT TIME OF SAMPLING																
55. REASON FOR SAMPLE: DYSMORPHIC FEATURES CONGENITAL ABNORMALITIES CONFIRMATION OF PRENATAL DIAGNOSIS OTHER																														
B. NEURAL TUBE DI																														
56. NEURAL TUBE DEFECT			ICD 9	Code	es 740	0.0-74	(2.0)						Т	57 S	PINA I	BIFIDA	Δ	FOR	All N	TD CAS	FS:				YES	3 1	10 UI	NKNOW	/N	
■ ANENCEPHALY			ACRAN	NIA		[SPIN	A BIFI	DA					YPE:						EPHALY PRESENT?									
RACHISCHISIS	11010			CEPHA		[A BIFII HYDF		PHALI	JS	OPEN 59. NT							PART OF A SYNDROME?										
CRANIORACHISCH	11515					5	Spina I	Bifida i	include	s Mer	ningon	yeloce	locele, CLOSED 60. IF YES.							SPECI	SPECIFY SYNDROME:									
INIENCEPHALY UNSPECIFIED Spile Initial Initiation with Initiat													61. ARE OTHE PRESENT								BNO	RMAI	LITIES	s					l	
MYELOMENINGOCELE														J UNS	SPEC	IFIED		Specify												
PRENATAL GENETIC PROCEDURES PERFORMED TO DETECT NEURAL TUBE DEFECT																														
PRENATAL DATE PROCEDURE PERFORMED NAME & ADDRESS OF FACE PROCEDURE (MM/DD/YYYY) WHERE PROCEDURE W																	ÞΕ					_ AGE KS/D				CEDU T NTD				
62. ULTRASOUND:		/		/																				/			YES	NO	I	
63. AMNIOCENTESIS:		/		/[/			YES	NO	I	
54. IF AMNIO PERFORMED, PROVIDE THE AF-AFP LEVEL (in M.o.M.) AT THE TIME OF LIVE BIRTH																	NOSE	D? 6	67. DATE OF POSTNATAL DIAGNOSIS:											
65. AF-AChE POSITIVE NEGATIVE AT P												YSICA	TIME OF LIVE BIRTH OTHER SICAL EXAMINATION Specify Other: E OF STILLBIRTH								1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1									
68. WAS THE FETAL ABNORMALITY CONFIRME] NO			F	PENDI	NG				□ NC	T PO	SSIBLE] UNI	KNOV	VN						
69. METHOD USED FOR AUTOPSY / PATHOLOGY VISUAL E																				OTHER Specify:										
ABNORMALITY: ULTRASOUND ONLY ULTRASO											UND & AMNIO									<u> </u>										
70. WHAT WAS THE SOURCE OF SOURCES OF YOUR CONFIRMATION? ULTRASOUND REPORT AMNIO RE DELIVERY DELIVERY ROOM RE											ERY	OUTCOME OTHER																		
PRIVACY The Infor	mation I	Dractices	. Act α	f 1077	(Civil (nde 1	708 0	t coa) requi	ree th	at the	followin	na dei	taile h	nrovi	ded w	hen a	form ie	used to	ohtair	info		•	m indiv	iduals	The	data re	nuesten	1 in	

PRIVACY STATEMENT The Information Practices Act of 1977 (Civil Code 1798 et. seq.) requires that the following details be provided when a form is used to obtain information from individuals. The data requested in this form are required by the Genetic Disease Branch (GDB) of the California Department of Health Services and are mandated by C.C.R.17 Section 6532. These data are used to provide information to subjects on the prevention of birth defects, to determine prevalence of chromosomal and neural tube defects and to monitor trends of occurrence. They will also be used to determine the effectiveness of the California Expanded Alpha Fetoprotein Screening Program. It is mandatory that health professionals completing this form provide complete and accurate information. The records maintained by the GDB are confidential as defined in Civil Code 1798.34 and are exempt from access by any individual except licensed medical personnel designated by the subject. The information may also be used in special studies as defined in Health and Safety Code 100330. The furnishing of such information to the Department or its authorized representative, or any other cooperating individual, agency or organization in any such special study shall not subject any person, hospital or other organization furnishing such information to any actions or damages.